Form PTO-1449 (REV. 7-80) U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

LIST OF PRIOR ART CITED BY APPLICANT

(Use several sheets if necessary)

Atty. Docket No.	Serial No.
18896	10/535,434
Applicants Kirby Siemering, et al.	
Filing Date September 14, 2006	**Group Art Unit** 1634

U.S. PATENT DOCUMENTS

EXAMINER INITIAL*		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE (if appropriate)

U.S. PATENT PUBLICATION DOCUMENTS

FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION	
							YES	NO

OTHER PRIOR ART (Including Author, Title, Date, Pertinent Pages, Etc.)

		Van Hauwe P. et al., "Two Frequent Missense Mutations in Pendred Syndrome", *Human Molecular Genetics, 7(7)*:1099-1104 (1998), XP-002454422
		Leroy B.P. et al., "Spectrum of Mutations in *USH2A* in British Patients with Usher Syndrome Type II", *Experimental Eye Research, 72(5)*:503-509 (2001), XP-002454423
		Nájera C. et al., "Mutations in Myosin VIIA (*MYO7A*) and Usherin (*USH2A*) in Spanish Patients with Usher Syndrome Types I and II, Respectively", *Human Mutation 20(1)*:1-7 (2002), XP-002454425
		Bogazzi F. et al., "A Novel Mutation in the Pendrin Gene Associated with Pendred's Syndrome", *Clinical Endocrinology, 52(3)*:279-285 (2000), XP-002454424
		Weston M.D. et al., "Genomic Structure and Identification of Novel Mutations in Usherin, the Gene Responsible for Usher Syndrome Type IIa", *American Journal of Human Genetics., 66(4)*:1199-1210 (2000), XP-002454426

EXAMINER	DATE CONSIDERED

*** EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.**